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Claims Amendment

Please add new claim 13.

1-8 (Cancelled)

9. (Previously submitted) A method for prophylaxis and /or treatment of endoparasitosis in an animal, comprising administration to the animal of an effective amount of a compound having the following formula (I):

wherein:

n is 0 or an integer comprised between 1 and 5;

R is a group R2-X -C(=Z)-NH-, in which X represents a simple chemical bond, an aromatic or heteroaromatic radical, Z represents an oxygen atom or the NH group; and:

if X is a simple chemical bond, R2 is an hydrogen atom, an alkyl,

dialkylaminoalkyl, alkenyl, cycloalkyl, arylalkyl, arylalkenyl, haloalkyl, or an aromatic or heteroaromatic radical;

if X is an aromatic or heteroaromatic radical, R2 is nitro, amino or formylamino;

or:

R is a group R3-C(=Z)-, in which Z represents an oxygen atom or the NH group, and R3 represents a hydrogen atom, the -OR4 or -NR5R6 group, and where:

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R4 is chosen from the group consisting of a hydrogen atom, an alkyl, cycloalkyl, arylalkyl, or an aromatic radical;

R5 and R6, either the same or different, are chosen from the group consisting of a hydrogen atom, an alkyl, cycloalkyl, arylalkyl, aromatic or heterocyclic radical, optionally substituted with a formylamino or a carbamoyl group; or

R5 and R6, joined together form an alkylene group, or the group -(CH2)2-O-(CH2)2- or the group -(CH2)2-NH-(CH2)2-;

A represents a simple chemical bond or the group -CO-NH-Y-, wherein Y is an alkylene or aromatic radical,

R1 is chosen from the group consisting of CH2N(CH3)2,-COOR4, -B-NR5R6, -C(=NH)-NH2, a heterocyclic radical containing nitrogen, wherein:

R4, R5 and R6 are as defined above, B represents a simple chemical bond or the -C=O group, and:

when R₁ is -B-NR₅R₆, and B is a simple chemical bond, or when R₁ is a heterocyclic radical, A is not a chemical bond;

or a pharmaceutical acceptable salt thereof.

10. (Previously submitted) The method of Claim 9, wherein the compound of formula (I) is chosen between distarrycin and a compound of formula (I) wherein:

n is as previously defined;

R is the -CONH2 group, A is the -CONHCH2CH2- group, R1 is the -C(⇒NH)-NH2 group or the -CH2N(CH3)2 group;

R is the -NH-CH(=NH) group, A is the -CONHCH2CH2- group, R₁ is the -C(=NH)-NH2 group or the -CH2N(CH3)2 group or the -CONH2 group; and the pharmaceutically acceptable salts thereof.

11. (Previously submitted) The method according to Claim 9, wherein the endoparasitosis is chosen from Trichomoniasis, Giardiasis, Istomoniasis, Amoebiasis, Coccidiosis, and Balantidiosis.

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- 12. (Previously submitted) The method according to Claim 9, wherein the administration is oral administration.
- 13. (New) The method according to Claim 1, wherein the compound of formula (I) is a compound of formula (I) wherein n is 0 or an integer from 1 to 5; R is a -CONH2 group; A is a -CONHCH2CH2- group, and R1 is a -C(=NH)-NH2 group, and the endoparasitosis is Coccidiosis.